IN THE CLAIMS

Please amend the claims as follows:

Claims 1-9 (Canceled).

Claim 10 (Currently Amended): A method for treatment of a Central Nervous System (CNS) disease, comprising administering, to a patient subject suffering from a disease of the CNS, a conjugate comprising an active substance in an amount sufficient to treat for treatment of said disease of the CNS, wherein the active substance is coupled directly or indirectly by a covalent bond to one of the following peptides: SynB1 (SEQ ID NO: 11) or SynB3 (SEQ ID NO: 12)[[;]] and treating said disease of the CNS, and wherein said active substance is an active chemical molecules molecule, and wherein said disease of the CNS is selected from the group consisting of brain cancer, pain and meningitis.

Claim 11 (Currently Amended): A method for driving a substance across the Blood Brain Barrier (BBB) to the Central Nervous System (CNS), comprising:

preparing a conjugate comprising an active substance coupled directly or indirectly by a covalent bond to one of the following peptides: SynB1 (SEQ ID NO: 11) or SynB3 (SEQ ID NO: 12), wherein said active substance is an active chemical molecules molecule in the treatment of the CNS;

administering said conjugate to a patient subject suffering from a disease of the CNS in an amount sufficient to drive said active substance across the BBB to the CNS; and driving one of the following peptides: SynB1 (SEQ ID NO: 11) or SynB3 (SEQ ID NO: 12) said active substance across the BBB to the CNS,

wherein said disease of the CNS is selected from the group consisting of brain cancer, pain, and meningitis.

Claim 12 (Previously Presented): The method of claim 10, wherein said active

chemical molecule is selected from the group consisting of antitumoral agents, antibiotic

agents and analgesic agents.

Claim 13 (Previously Presented): The method of claim 11, wherein said active

chemical molecule is selected from the group consisting of antitumoral agents, antibiotic

agents and analgesic agents.

Claim 14 (New): The method of claim 10, wherein the amount of said conjugate

administered to the subject is from 0.16 to 2.5 mg/kg of body weight.

Claim 15 (New): The method of claim 11, wherein the amount of said conjugate

administered to the subject is from 0.3 to 3 mg/kg of body weight.

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